Yasue Horiuchi

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A novel susceptibility locus for moyamoya disease on chromosome 8q23. Journal of Human Genetics, 2004, 49, 278-281.	1.1	187
2	Brain Cannabinoid CB2 Receptor in Schizophrenia. Biological Psychiatry, 2010, 67, 974-982.	0.7	163
3	Involvement of cannabinoid CB2 receptor in alcohol preference in mice and alcoholism in humans. Pharmacogenomics Journal, 2007, 7, 380-385.	0.9	119
4	Involvement of SMARCA2/BRM in the SWI/SNF chromatin-remodeling complex in schizophrenia. Human Molecular Genetics, 2009, 18, 2483-2494.	1.4	103
5	Pathway-based association analysis of genome-wide screening data suggest that genes associated with the γ-aminobutyric acid receptor signaling pathway are involved in neuroleptic-induced, treatment-resistant tardive dyskinesia. Pharmacogenetics and Genomics, 2008, 18, 317-323.	0.7	95
6	Deficits in microRNA-mediated Cxcr4/Cxcl12 signaling in neurodevelopmental deficits in a 22q11 deletion syndrome mouse model. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 17552-17557.	3.3	65
7	A polymorphism of the metabotropic glutamate receptor mGluR7 (GRM7) gene is associated with schizophrenia. Schizophrenia Research, 2008, 101, 9-16.	1.1	59
8	A nonsynonymous polymorphism in cannabinoid CB2 receptor gene is associated with eating disorders in humans and food intake is modified in mice by its ligands. Synapse, 2010, 64, 92-96.	0.6	57
9	Association of the HSPG2 Gene with Neuroleptic-Induced Tardive Dyskinesia. Neuropsychopharmacology, 2010, 35, 1155-1164.	2.8	57
10	Olfactory cells via nasal biopsy reflect the developing brain in gene expression profiles: Utility and limitation of the surrogate tissues in research for brain disorders. Neuroscience Research, 2013, 77, 247-250.	1.0	51
11	Generation of Induced Pluripotent Stem Cells from Human Nasal Epithelial Cells Using a Sendai Virus Vector. PLoS ONE, 2012, 7, e42855.	1.1	46
12	Possible association between a haplotype of the GABA-A receptor alpha 1 subunit gene (GABRA1) and mood disorders. Biological Psychiatry, 2004, 55, 40-45.	0.7	40
13	Pyridoxamine: A novel treatment for schizophrenia with enhanced carbonyl stress. Psychiatry and Clinical Neurosciences, 2018, 72, 35-44.	1.0	40
14	Support for association of the PPP3CC gene with schizophrenia. Molecular Psychiatry, 2007, 12, 891-893.	4.1	38
15	DPP6 as a candidate gene for neuroleptic-induced tardive dyskinesia. Pharmacogenomics Journal, 2013, 13, 27-34.	0.9	38
16	A Polymorphism in the PDLIM5 Gene Associated with Gene Expression and Schizophrenia. Biological Psychiatry, 2006, 59, 434-439.	0.7	37
17	Association of polymorphisms in the haplotype block spanning the alternatively spliced exons of the NTNC1 gene at 1p13.3 with schizophrenia in Japanese populations. Neuroscience Letters, 2008, 435, 194-197.	1.0	37
18	Monoallelic and Unequal Allelic Expression of the HTR2A Gene in Human Brain and Peripheral Lymphocytes. Biological Psychiatry, 2006, 60, 1331-1335.	0.7	36

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19	Functional polymorphism in the <i>GPR55</i> gene is associated with anorexia nervosa. Synapse, 2011, 65, 103-108.	0.6	36
20	Valley of death: A proposal to build a "translational bridge―for the next generation. Neuroscience Research, 2017, 115, 1-4.	1.0	33
21	New Insights and Potential Therapeutic Targeting of CB2 Cannabinoid Receptors in CNS Disorders. International Journal of Molecular Sciences, 2022, 23, 975.	1.8	32
22	RGS4 is not a susceptibility gene for schizophrenia in Japanese: Association study in a large case-control population. Schizophrenia Research, 2007, 89, 161-164.	1.1	30
23	Molecular signatures associated with cognitive deficits in schizophrenia: a study of biopsied olfactory neural epithelium. Translational Psychiatry, 2016, 6, e915-e915.	2.4	30
24	Carbonyl stress and schizophrenia. Psychiatry and Clinical Neurosciences, 2014, 68, 655-665.	1.0	29
25	<scp>NrCAM</scp> â€regulating neural systems and addictionâ€related behaviors. Addiction Biology, 2014, 19, 343-353.	1.4	29
26	Cannabinoid CB2 Receptor Gene and Environmental Interaction in the Development of Psychiatric Disorders. Molecules, 2018, 23, 1836.	1.7	28
27	Failure to find causal mutations in the GABAA-receptor γ2 subunit (GABRG2) gene in Japanese febrile seizure patients. Neuroscience Letters, 2003, 343, 117-120.	1.0	25
28	Association of SNPs linked to increased expression of SLC1A1 with schizophrenia. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 30-37.	1.1	25
29	Determination of methylglyoxal in human blood plasma using fluorescence high performance liquid chromatography after derivatization with 1,2-diamino-4,5-methylenedioxybenzene. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2016, 1029-1030, 102-105.	1.2	24
30	Supportive Evidence for Reduced Expression of GNB1L in Schizophrenia. Schizophrenia Bulletin, 2010, 36, 756-765.	2.3	23
31	Replication study and meta-analysis of the genetic association of GRM3 gene polymorphisms with schizophrenia in a large Japanese case-control population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 392-396.	1.1	20
32	Experimental Evidence for the Involvement of PDLIM5 in Mood Disorders in Hetero Knockout Mice. PLoS ONE, 2013, 8, e59320.	1.1	18
33	A 58-kDa Shc Protein Is Present inXenopusEggs and Is Phosphorylated on Tyrosine Residues upon Egg Activation. Biochemical and Biophysical Research Communications, 1999, 258, 265-270.	1.0	17
34	Failure to confirm the association between the FEZ1 gene and schizophrenia in a Japanese population. Neuroscience Letters, 2007, 417, 326-329.	1.0	16
35	From population to neuron: exploring common mediators for metabolic problems and mental illnesses. Molecular Psychiatry, 2021, 26, 3931-3942.	4.1	16
36	Disclosure of secondary findings in exome sequencing of 2480 Japanese cancer patients. Human Genetics, 2021, 140, 321-331.	1.8	16

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37	Vitamin B6 deficiency hyperactivates the noradrenergic system, leading to social deficits and cognitive impairment. Translational Psychiatry, 2021, 11, 262.	2.4	16
38	Identification of an argpyrimidine-modified protein in human red blood cells from schizophrenic patients: A possible biomarker for diseases involving carbonyl stress. Biochemical and Biophysical Research Communications, 2017, 493, 573-577.	1.0	15
39	Replication study of association between ADCYAP1 gene polymorphisms and schizophrenia. Psychiatric Genetics, 2010, 20, 123-125.	0.6	15
40	Characterization of modified proteins in plasma from a subtype of schizophrenia based on carbonyl stress: Protein carbonyl is a possible biomarker of psychiatric disorders. Biochemical and Biophysical Research Communications, 2015, 467, 361-366.	1.0	14
41	The regulation of soluble receptor for AGEs contributes to carbonyl stress in schizophrenia. Biochemical and Biophysical Research Communications, 2016, 479, 447-452.	1.0	14
42	Clinical Utility of Neuronal Cells Directly Converted from Fibroblasts of Patients for Neuropsychiatric Disorders: Studies of Lysosomal Storage Diseases and Channelopathy. Current Molecular Medicine, 2015, 15, 138-145.	0.6	14
43	Potential Role of Cannabinoid Type 2 Receptors in Neuropsychiatric and Neurodegenerative Disorders. Frontiers in Psychiatry, 0, 13, .	1.3	14
44	PICK1 is not a susceptibility gene for schizophrenia in a Japanese population: Association study in a large case–control population. Neuroscience Research, 2007, 58, 145-148.	1.0	13
45	Combined glyoxalase 1 dysfunction and vitamin B6 deficiency in a schizophrenia model system causes mitochondrial dysfunction in the prefrontal cortex. Redox Biology, 2021, 45, 102057.	3.9	12
46	Replication study for associations between polymorphisms in the CLDN5 and DGCR2 genes in the 22q11 deletion syndrome region and schizophrenia. Psychiatric Genetics, 2008, 18, 255-256.	0.6	11
47	Germline mismatch repair gene variants analyzed by universal sequencing in Japanese cancer patients. Cancer Medicine, 2019, 8, 5534-5543.	1.3	10
48	Role of glyoxalase 1 in methylglyoxal detoxification–the broad player of psychiatric disorders. Redox Biology, 2022, 49, 102222.	3.9	9
49	Prevalence of low-penetrant germline <i>TP53</i> D49H mutation in Japanese cancer patients . Biomedical Research, 2016, 37, 259-264.	0.3	8
50	Role of Tumor Mutation Burden Analysis in Detecting Lynch Syndrome in Precision Medicine: Analysis of 2,501 Japanese Cancer Patients. Cancer Epidemiology Biomarkers and Prevention, 2021, 30, 166-174.	1.1	8
51	Dysregulation of post-transcriptional modification by copy number variable microRNAs in schizophrenia with enhanced glycation stress. Translational Psychiatry, 2021, 11, 331.	2.4	7
52	Advanced glycation end products and cognitive impairment in schizophrenia. PLoS ONE, 2021, 16, e0251283.	1.1	6
53	Fingertip advanced glycation end products and psychotic symptoms among adolescents. NPJ Schizophrenia, 2021, 7, 37.	2.0	6
54	Microsatellite instability is biased in Amsterdam II-defined Lynch-related cancer cases with family history but is rare in other cancers: a summary of 1000 analyses. BMC Cancer, 2022, 22, 73.	1.1	5

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55	A case of type 1 multiple endocrine neoplasia with esophageal stricture successfully treated with endoscopic balloon dilation and local steroid injection combined with surgical resection of gastrinomas. BMC Gastroenterology, 2017, 17, 37.	0.8	4
56	Metachronous ovarian endometrioid carcinomas in a patient with a PTEN variant: case report of incidentally detected Cowden syndrome. BMC Cancer, 2019, 19, 1014.	1.1	3
57	Cooperation of LIM domainâ€binding 2 (LDB2) with EGR in the pathogenesis of schizophrenia. EMBO Molecular Medicine, 2021, 13, e12574.	3.3	2
58	Present status of germline findings in precision medicine for Japanese cancer patients: issues in the current system. Japanese Journal of Clinical Oncology, 2022, 52, 599-608.	0.6	2
59	Germline and somatic genetic changes in multicentric tumors obtained from a patient with multiple endocrine neoplasia type 1. Human Genome Variation, 2017, 4, 17013.	0.4	1
60	A novel MLH1 intronic variant in a young Japanese patient with Lynch syndrome. Human Genome Variation, 2018, 5, 3.	0.4	1
61	LDB2 locus disruption on 4p16.1 as a risk factor for schizophrenia and bipolar disorder. Human Genome Variation, 2020, 7, 31.	0.4	1
62	Exonic deletions in IMMP2L in schizophrenia with enhanced glycation stress subtype. PLoS ONE, 2022, 17, e0270506.	1.1	1
63	Generation of induced pluripotent stem cells from patients with schizophrenia. Neuroscience Research, 2010, 68, e314.	1.0	0
64	Fabry disease has been found by using of the tumor mutational burden analysis of 3000 Japanese cancer genomes using whole exome and targeted gene panel sequencing: Project Hightech Omics-based Patient Evaluation (Project HOPE). Molecular Genetics and Metabolism, 2019, 126, S154-S155.	0.5	0
65	Genomic profiling of multiple tissues in two patients with multiple endocrine neoplasia type 1. Biomedical Research, 2021, 42, 89-94.	0.3	0